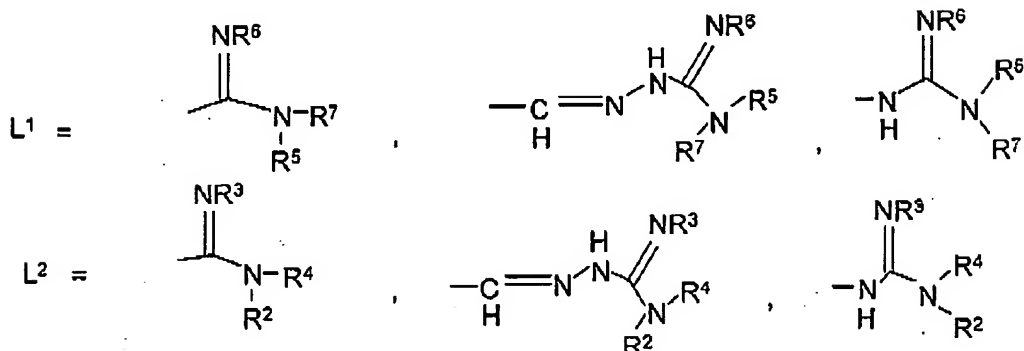
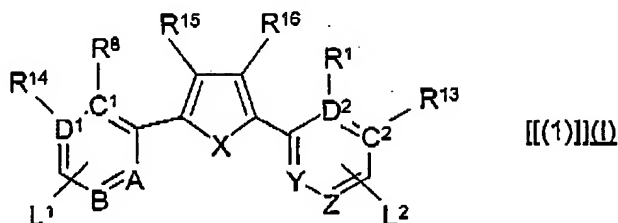


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IN THE CLAIMS:

Please amend the claims as follows:

1. (Currently amended) A compound of Formula (I):



wherein:

X is selected from the group consisting of O, S, and NR^{17} , where R^{17} is hydrogen or lower alkyl;

C^1 , C^2 , A, Y, and Z are CH, N, NR^{17} , O, or S;

C^1 and C^2 are each C or N, wherein C^1 and C^2 are the same or different;

D^1 and D^2 are each CH, N, or NR^{17} , wherein D^1 and D^2 are the same or different;

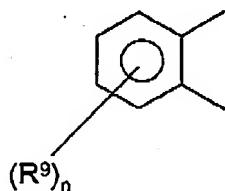
B and Z are CH, N, or NR^{17} , provided that B, Z, or both B and Z are not present when A, Y, or both A and Y are O, S, or NR^{17} ;

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R^{13} , R^{14} , R^{15} , R^{16} , R^1 and R^8 can be present or absent, and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

R^{15} and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

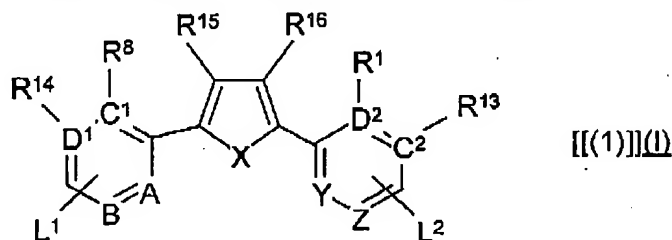
R^3 and R^6 are each independently selected from the group consisting of H, hydroxy, lower alkyl, cycloalkyl, aryl, aralkyl, alkoxy, hydroxycycloalkyl, alkoxyalkyl, hydroxyalkyl, aminoalkyl, acyloxy, acetoxy, and alkylaminoalkyl; and R^2 , R^4 , R^5 and R^7 are each independently selected from the group consisting of H, lower alkyl, alkoxyalkyl, cycloalkyl, aryl, aralkyl, hydroxyalkyl, aminoalkyl, and alkylaminoalkyl, or R^2 and R^4 together or R^5 and R^7 together represent a C_2 to C_{10} alkyl, hydroxyalkyl, or alkylene, or R^3 and R^4 together or R^6 and R^7 together are:



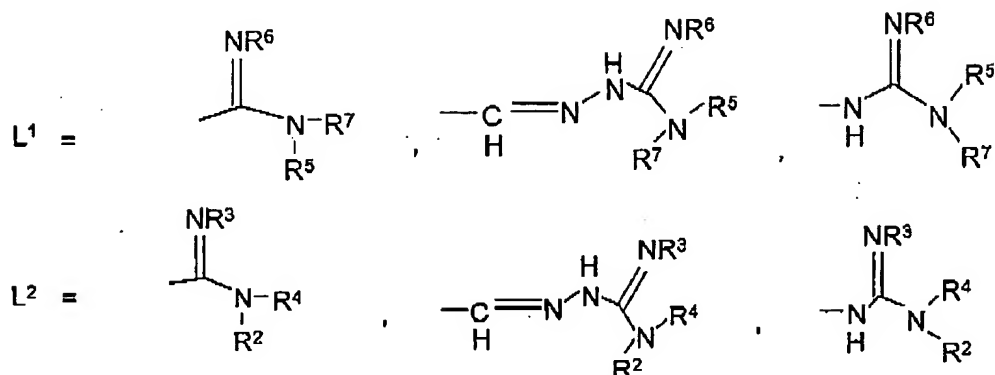
wherein n is a number from 1 to 3, and R^9 is H or $-\text{CONHR}^{10}\text{NR}^{11}\text{R}^{12}$, wherein R^{10} is lower alkyl and R^{11} and R^{12} are each independently selected from the group consisting of H and lower alkyl; and

wherein at least one of A, B, Y, and Z are selected from the group consisting of N, NR^{17} , O, and S.

2. (Currently amended) A compound of Formula (I):

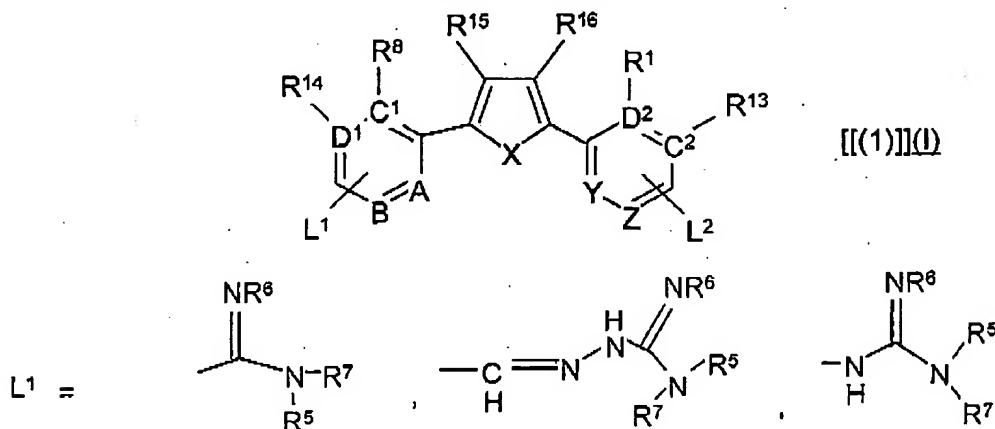


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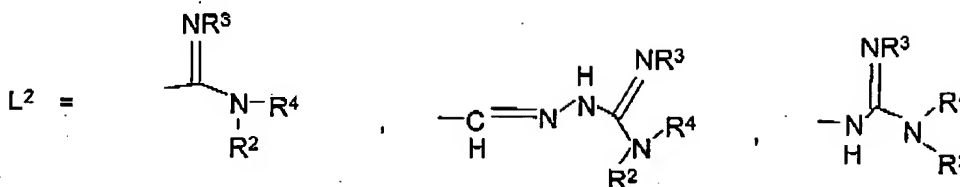


wherein A and B are different and N or CH; Y and Z are CH; X is O or S; R^2 , R^4 , R^5 , and R^7 are each H; $[[R^1,]]$ $R^3[[,]]$ and R^6 and R^8 are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R^1 and R^8 can be present or absent and when present are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; $R^{13}[[,]]$ and R^{14} can be present or absent and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; $[[,]]$ $R^{15}[[,]]$ and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; and C^1 , C^2 , D^1 , and D^2 are each C[[H]] or N.

3. (Currently amended) A compound of Formula (I):



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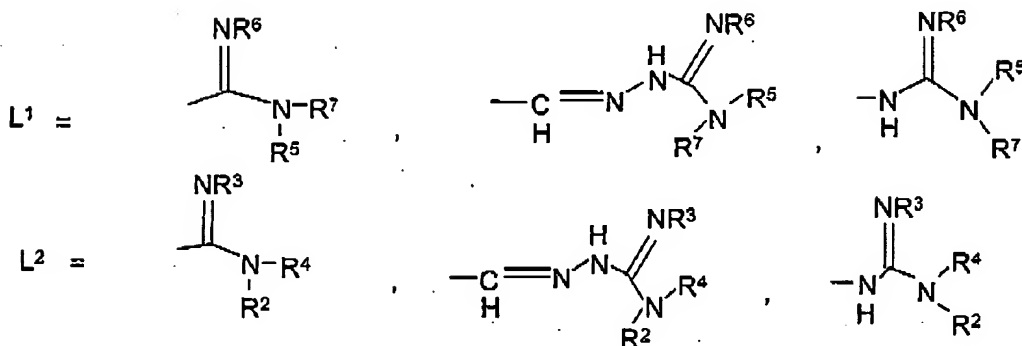
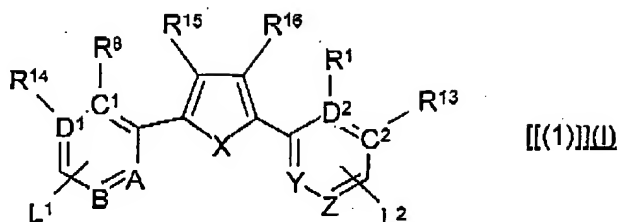


wherein A and B are CH; X is O; Y is O; Z is not present; R^2 , R^4 , R^5 , and R^7 are each H; $[[R^1,]]$ $R^3[[,]]$ and R^6 and R^8 are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R^1 and R^8 can be present or absent and when present are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; $R^{13}[[,]]$ and R^{14} can be present or absent and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; $[[,]]$ $R^{15}[[,]]$ and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; and C^1 , C^2 , D^1 , and D^2 are each C[[H]] or N.

4. (Original) The compound of claim 1, further comprising a pharmaceutically acceptable carrier.

5. (Currently amended) A method of treating a microbial infection in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I):

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wherein:

X is selected from the group consisting of O, S, and NR^{17} , where R^{17} is hydrogen or lower alkyl;

C^1 , C^2 , A, B, and Y are CH, N, NR^{17} , O, or S;

C^1 and C^2 are each C or N, wherein C^1 and C^2 are the same or different;

D^1 , D^2 , B, and Z are CH, N, or NR^{17} , wherein D^1 and D^2 are the same or different;

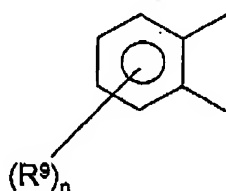
B and Z are CH, N, or NR^{17} , provided that B, Z, or both B and Z are not present when A, Y, or both A and Y are O, S, or NR^{17} ;

R^{13} , R^{14} , R^{15} , R^{16} , R^1 and R^8 can be present or absent, and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

R^{15} and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

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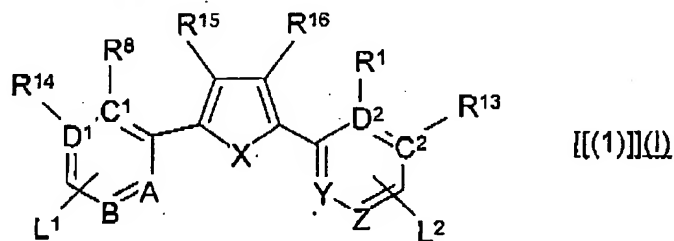
R^3 and R^6 are each independently selected from the group consisting of H, hydroxy, lower alkyl, cycloalkyl, aryl, aralkyl, alkoxy, hydroxycycloalkyl, alkoxyalkyl, hydroxyalkyl, aminoalkyl, acyloxy, acetoxy, and alkylaminoalkyl; and R^2 , R^4 , R^5 and R^7 are each independently selected from the group consisting of H, lower alkyl, alkoxyalkyl, cycloalkyl, aryl, aralkyl, hydroxyalkyl, aminoalkyl, and alkylaminoalkyl, or R^2 and R^4 together or R^5 and R^7 together represent a C_2 to C_{10} alkyl, hydroxyalkyl, or alkylene, or R^3 and R^4 together or R^6 and R^7 together are:



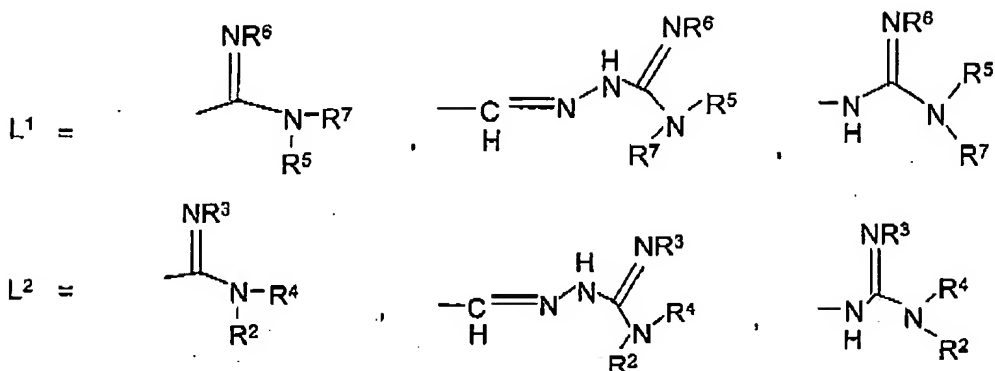
wherein n is a number from 1 to 3, and R^9 is H or $-\text{CONHR}^{10}\text{NR}^{11}\text{R}^{12}$, wherein R^{10} is lower alkyl and R^{11} and R^{12} are each independently selected from the group consisting of H and lower alkyl; and

wherein at least one of A, B, Y, and Z are selected from the group consisting of N, NR^{17} , O, and S.

6. (Currently amended) A method of treating a microbial infection in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I):

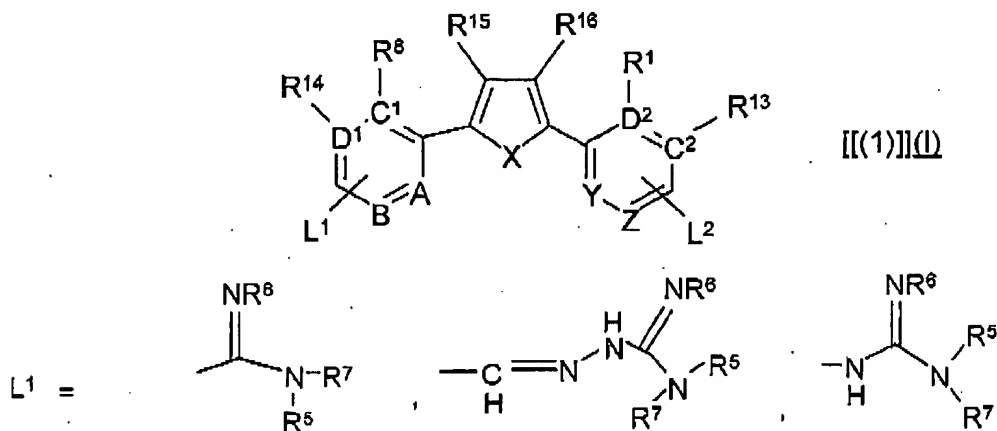


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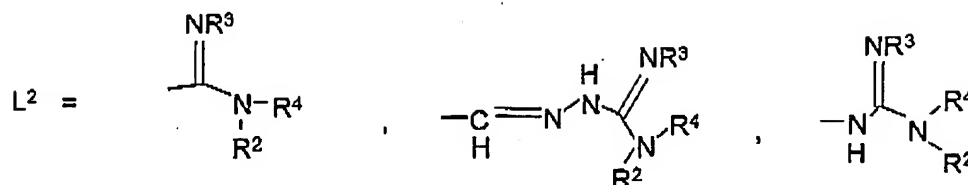


wherein A and B are different and N or CH; Y and Z are CH; X is O or S; R², R⁴, R⁵, and R⁷ are each H; [[R¹,]] R³[[,]] and R⁶ and R⁸ are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R¹ and R⁸ can be present or absent and when present are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R¹³[[,]] and R¹⁴ can be present or absent and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;[[,]] R¹⁵[[,]] and R¹⁶ are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; and C¹, C², D¹, and D² are each C[[H]] or N.

7. (Currently amended) A method of treating a microbial infection in a subject in need thereof, the method comprising administering to the subject an effective amount of a compound of Formula (I):



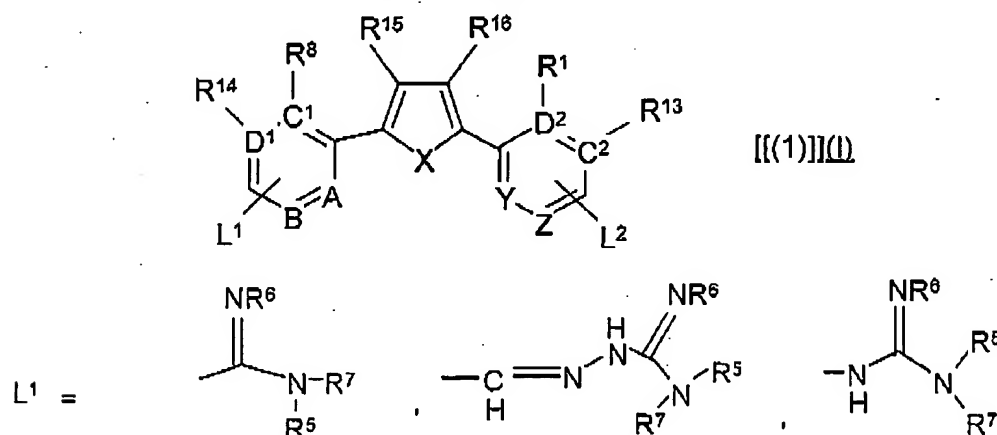
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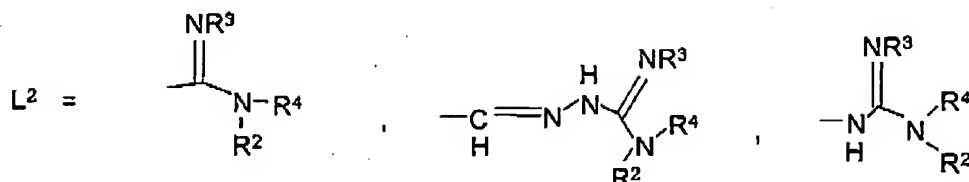
wherein A and B are CH; X is O; Y is O; Z is not present; R², R⁴, R⁵, and R⁷ are each H; [R¹,] R³[.,] and R⁶ and ~~R⁸~~ are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R¹ and R⁸ can be present or absent and when present are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R¹³[.,] and R¹⁴ can be present or absent and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;[.,] R¹⁵[.,] and R¹⁶ are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; and C¹, C², D¹, and D² are each C[H] or N.

8. (Original) The method of claim 5, wherein the microbial infection is a *Trypanosoma brucei rhodesiense* infection or a *Plasmodium falciparum* infection.

9. (Currently amended) A pharmaceutical formulation comprising:
(a) a compound of Formula (I):



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wherein:

X is selected from the group consisting of O, S, and NR^{17} , where R^{17} is hydrogen or lower alkyl;

$\text{G}^1, \text{G}^2, \text{A}[[.]]$ and Y are CH, N, NR^{17} , O, or S;

C^1 and C^2 are each C or N, wherein C^1 and C^2 are the same or different;

$\text{D}^1[[.]]$ and D^2, B , and Z are CH, N, or NR^{17} are each C or N, wherein D^1 and D^2 are the same or different;

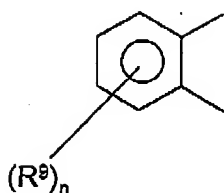
B and Z are CH, N, or NR^{17} , provided that B, Z, or both B and Z are not present when A, Y, or both A and Y are O, S, or NR^{17} ;

$\text{R}^{13}, \text{R}^{14}, \text{R}^{15}, \text{R}^{16}, \text{R}^1$ and R^8 can be present or absent, and when present are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

R^{15} and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl;

R^3 and R^6 are each independently selected from the group consisting of H, hydroxy, lower alkyl, cycloalkyl, aryl, aralkyl, alkoxy, hydroxycycloalkyl, alkoxy-cycloalkyl, hydroxyalkyl, aminoalkyl, acyloxy, acetoxy, and alkylaminoalkyl; and $\text{R}^2, \text{R}^4, \text{R}^5$ and R^7 are each independently selected from the group consisting of H, lower alkyl, alkoxyalkyl, cycloalkyl, aryl, aralkyl, hydroxyalkyl, aminoalkyl, and alkylaminoalkyl, or R^2 and R^4 together or R^5 and R^7 together represent a C_2 to C_{10} alkyl, hydroxyalkyl, or alkylene, or R^3 and R^4 together or R^6 and R^7 together are:

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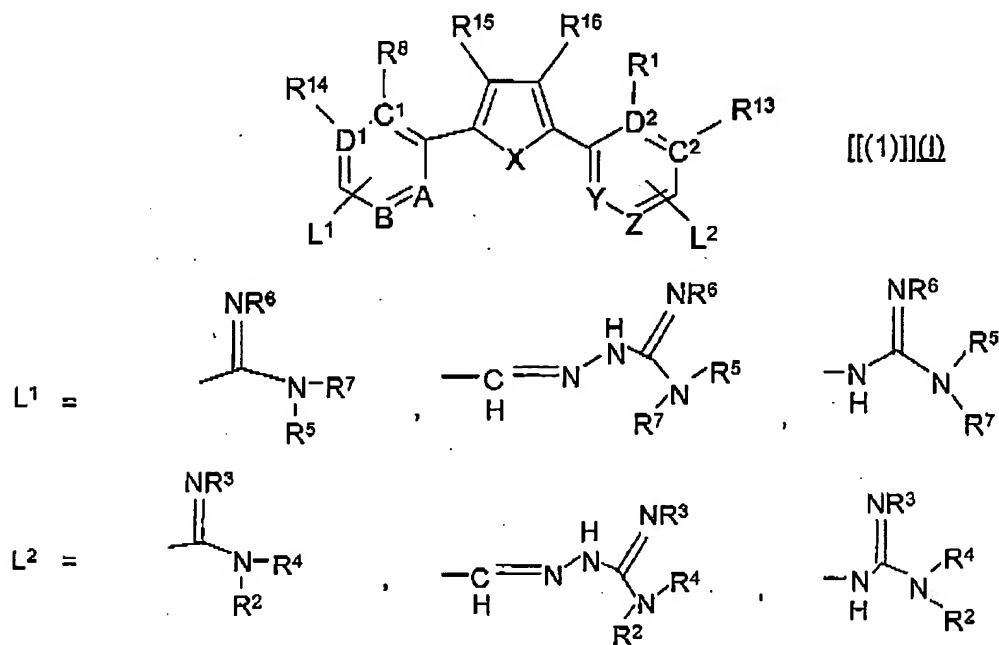
wherein n is a number from 1 to 3, and R^9 is H or $-\text{CONHR}^{10}\text{NR}^{11}\text{R}^{12}$, wherein R^{10} is lower alkyl and R^{11} and R^{12} are each independently selected from the group consisting of H and lower alkyl; and

wherein at least one of A, B, Y, and Z are selected from the group consisting of N, NR^{17} , O, and S; and

(b) a pharmaceutically acceptable carrier.

10. (Currently amended) A pharmaceutical formulation comprising:

(a) a compound of Formula (I):



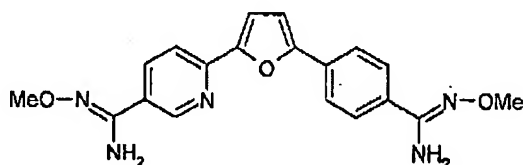
wherein A and B are different and N or CH; Y and Z are CH; X is O or S; R^2 , R^4 , R^5 , and R^7 are each H; $[(R^1)]$, R^3 , $[(L)]$ and R^6 and R^8 are selected from the group consisting of H, OH, methyl, methoxy, and acetoxy; R^1

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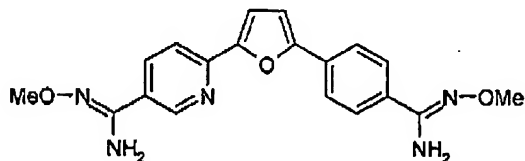
consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; [[,]] R^{15} [[,]] and R^{16} are selected from the group consisting of H, lower alkyl, halogen, alkoxy, aryloxy, aralkoxy and hydroxyl; and C^1 , C^2 , D^1 , and D^2 are each C[[H]] or N; and

(b) a pharmaceutically acceptable carrier.

12. (Previously presented) The compound of claim 2, wherein A is N; B is CH; X is O; R_1 and R_8 are H; R_3 and R_6 are methoxy; and the compound has the structure:



13. (Previously presented) The method of claim 6, wherein A is N; B is CH; X is O; R_1 and R_8 are H; R_3 and R_6 are methoxy; and the compound has the structure:



14. (Previously presented) The method of claim 13, wherein the microbial infection is a *Trypanosoma brucei rhodesiense* infection or a *Plasmodium falciparum* infection.

15. (Previously presented) The pharmaceutical formulation of claim 10, wherein A is N; B is CH; X is O; R_1 and R_8 are H; R_3 and R_6 are methoxy; and the compound has the structure:

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